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1-13. (canceled)

14. A method for producing *Vesicular stomatitis* virus G envelope protein (VSV-G) pseudotyped vector particles or virus like particles thereof containing a transgene, the method comprising:

obtaining cells from a packaging cell line that is negative for expression of low density lipoprotein receptor (LDLR);

causing the cells to express VSV-G, gag/pol, and optionally rev; and

causing the cells to express the transgene;

wherein expression of the aforesaid genes in the cells results in production of the vector particles or virus-like particles containing the transgene; then

harvesting from the cells the vector particles or virus-like particles.

15. The method of claim 14, wherein the packaging cell line is a human cell line.

16. The method of claim 14, wherein the packaging cell line has been genetically engineered to prevent expression of LDLR on the cell surface.

17. The method of claim 16, wherein lentivirus autotransduction by the packaging cell line is reduced by at least 50% as a result of being genetically engineered to prevent expression of LDLR on the cell surface.

18. The method of claim 14, wherein the packaging cell line is selected from human embryonic kidney (HEK) cell line 293, HEK 293T, HEK EBNA, HEK 293F, HEK 293FT, and HEK 293-S.

19. The method of claim 14, wherein the packaging cell line has been genetically engineered to stably express VSV-G.

20. The method of claim 19, wherein the packaging cell line has been genetically engineered to stably express at least one additional gene selected from Gag/pol, rev, and the transgene.

21. The method of claim 14, wherein the cells are caused to express VSV-G, gag/pol, and the transgene by a process that includes introducing into the cells:

a first Ψ -negative expression vector that encodes gag/pol, a second Ψ -negative expression vector that encodes rev if the vector particles or virus like particles being produced constitute a lentiviral expression vector,

a third Ψ -negative expression vector that encodes VSV-G, and

a Ψ -positive expression vector that contains the transgene.

22. The method of claim 21, wherein the cells are caused to express VSV-G, gag/pol, and the transgene by a process that includes co-transfecting the cells with:

a first Ψ -negative retroviral expression vector that encodes gag/pol,

a second Ψ -negative expression vector that encodes rev if the vector particles or virus like particles being produced constitute a lentiviral expression vector,

a third Ψ -negative retroviral expression vector that encodes VSV-G, and

a Ψ -positive retroviral expression vector that contains the transgene.

23. The method of claim 14, wherein the cells are caused to express VSV-G, gag/pol, rev, and the transgene by a process that includes co-transfecting the cells with:

a first Ψ -negative retroviral expression vector that encodes gag/pol/rev,

a second Ψ -negative retroviral expression vector that encodes VSV-G, and

a Ψ -positive retroviral expression vector that contains the transgene.

24. The method of claim 19, wherein the cells that stably express VSV-G are caused to express gag/pol and the transgene by a process that includes introducing into the cells:

a first Ψ -negative expression vector that encodes gag/pol,

a second Ψ -negative expression vector that encodes rev if the vector particles or virus like particles being produced constitute a lentiviral expression vector,

a Ψ -positive expression vector that contains the transgene.

25. The method of claim 20, wherein the cells are caused to express gag/pol, rev, and the transgene by a process that includes introducing into the cells:

Ψ -negative expression vector(s) that encode genes selected from gag/pol, rev, and VSV-G that are not constitutively expressed by the packaging cell line; and

a Ψ -positive expression vector contain the transgene.

26. The method of claim 14, further comprising formulating the harvested vector particles or virus-like particles as a pharmaceutical composition in a pharmaceutically acceptable carrier.

27. A method of treating a subject in need thereof, comprising administering to the subject a pharmaceutical composition produced according to the method of claim 26.

28. A combination configured for producing vesicular stomatitis virus G envelope protein (VSV-G) pseudotyped vector particles or virus like particles thereof containing a transgene, the combination comprising:

cells from a packaging cell line that is negative for expression of low density lipoprotein receptor (LDLR); along with

retrovirus expression vector(s) that encode one or more genes selected from VSV-G, gag/pol, and rev.